AMENDMENTS TO THE CLAIMS:

Claims 1-5 (Canceled)

6. (Currently Amended) A transgenic mouse emprising whose genome comprises a homozygous disruption in the FPR-RS4 gene, wherein the transgenic mouse exhibits, relative to a wild-type mouse, a phenotypic abnormality selected from the group consisting of increased anxiety, decreased coordination, impaired balance and decreased susceptibility to seizure.

Claim 7 (Canceled)

- 8. (Previously Presented) A cell derived from the transgenic mouse of claim 6.
- 9. (Currently Amended) A method of producing a transgenic mouse comprising whose genome comprises a homozygous disruption in the FPR-RS4 gene, the method comprising:
 - (a) introducing a construct that targets the FPR-RS4 gene into a mouse embryonic stem cell;
 - (b) introducing the embryonic stem cell into a blastocyst;
 - (c) implanting the resulting blastocyst into a pseudopregnant mouse, wherein said pseudopregnant mouse gives birth to a chimeric mouse; and
 - (d) breeding the chimeric mouse to produce the transgenic mouse, wherein the transgenic mouse exhibits, relative to a wild-type mouse, a phenotypic abnormality selected from the group consisting of increased anxiety, decreased coordination and decreased susceptibility to seizure.

Claims 10-22 (Canceled)

- 23. (Currently Amended) A method of identifying an agent that ameliorates a phenotype associated with a homozygous disruption in <u>the FPR-RS4 gene</u>, the method comprising:
 - (a) administering an agent to a transgenic mouse comprising a homozygous disruption in the-FPR-RS4gene; and
 - (b) determining whether the agent ameliorates at least one of the following phenotypes: increased anxiety, impaired motor coordination or balance, ataxia, or decreased susceptibility to seizure.

Claims 24-28 (Canceled)

29. (Previously Presented) A method of identifying an agent that ameliorates anxiety, the method comprising:

- (a) administering an agent to the transgenic mouse of claim-6; and
- (b) determining whether the agent has an affect on anxiety in the transgenic mouse.
- 30. (Currently Amended) A method of identifying an agent that ameliorates impaired motor coordination, impaired balance, or ataxia, the method comprising:
 - (a) administering an agent to the transgenic mouse of claim 6; and
 - (b) determining whether the agent has an affect on motor coordination, or balance-or ataxia in the transgenic mouse.
- 31. (Currently Amended) A method of evaluating treatments for anxiety, the method comprising:
 - (a) administering a therapeutic agent to the transgenic mouse of claim 6; and
 - (b) determining the *in vivo* effects of whether the agent has an effect on anxiety level in the transgenic mouse.
- 32. (Currently Amended) A method of evaluating treatments for impaired motor coordination or, impaired balance, or ataxia, the method comprising:
 - (a) administering a therapeutic agent to the transgenic mouse of claim 6; and
 - (b) determining the *in vivo* effects of whether the agent has an effect on motor coordination or, balance, or ataxia in the transgenic mouse.

Claims 33-34 (Canceled)

- 35. (Previously Presented) The transgenic mouse of claim 6, wherein the increased anxiety is characterized by decreased time spent in a central region during an open field test.
- 36. (Previously Presented) The transgenic mouse of claim 6, wherein the decreased coordination is characterized by decreased time to fall during a rotarod test.
- 37. (Currently Amended) The transgenic mouse of claim 6, wherein the decreased coordination is characterized by a decrease in time to fall falling off the accelerating rotarod at a lower speed.
- 38. (Previously Presented) The transgenic mouse of claim 6, wherein the decreased coordination comprises impaired motor coordination, impaired balance, or ataxia.
- 39. (Previously Presented) The transgenic mouse of claim 6, wherein the decreased susceptibility to seizure is characterized by an increased dose of metrazol to reach seizure.